CLAIMS

1. An intravitreous injectable solution for the treatment of vitreous hemorrhages, which comprises: a pharmaceutically effective quantity of an active ingredient; and a pharmaceutically acceptable quantity of a carrier solution.

2. The intravitreous injectable solution, according to claim 1, characterized in that said active ingredient is mannitol.

3. The intravitreous injectable solution, according to claim 1, characterized in that said carrier solution comprises: 10.20% in weight of polyoxyl stearate 40; 15% in weight of edetate disodium, dihydrate; 1.03% in weight of sodium chloride; 0.14% in weight of boric acid; 0.32% in weight of sorbic acid; 0.06% in weight of sodium bisulfate and 88.00% in weight of distilled water.

4. The intravitreous injectable solution, according to claim 1, characterized in that said carrier solution is Sophisen.

5. The intravitreous injectable solution, according to claim 1, characterized in that mannitol is present in a concentration that encourages the reabsorption of the vitreous hemorrhage.

6. The intravitreous injectable solution, according to claim 3, characterized in that the mannitol is present in the solution in a percentage of 5% to 30% in weight.

7. The intravitreous injectable solution, according to claim 1, characterized in that the Sophisen is present in the solution in a percentage of 0.02% to 20% in weight.

8. The intravitreous injectable solution, according to claim 1, characterized in that the pH of the solution is approximately 7.2.

9. The intravitreous injectable solution, according to claim 1, characterized in that the solution has an osmolarity of approximately 1400

mOsm/kg.

10. A method for the treatment of vitreous hemorrhages comprising: applying at least one injection of a pharmaceutically acceptable ophthalmic solution, into the vitreous humor of an eye with a hemorrhage resulting from a lesion or disease.

11. The method according to claim 10, characterized by applying at least one therapeutically effective dose of the injectable solution of claim 1, into the vitreous humor of a patient diagnosed with vitreous hemorrhage.

12. A method for the clarification of vitreous hemorrhages comprising the injection, into the vitreous body of the eye of a patient, of an ophthalmic solution, such as the one claimed in claim 1.

13. The method of the preceding claim, which is aimed at avoiding vitrectomy surgery in patients with vitreous hemorrhage, by the application of at least one intraocular injection of an ophthalmic solution formulated for the reabsorption of the hemorrhage.

14. The new use of mannitol as active ingredient of a solution that is injectable into the vitreous body of the eye for the treatment of vitreous hemorrhages.

15. A method for the preparation of an ophthalmic solution for an intravitreous injection for the treatment of vitreous hemorrhages, characterized by the following steps: pouring into a stainless steel recipient 800 ml of injectable water at a temperature of 40° ± 2°C; beginning the agitation at 200 rpm ± 50 rpm and keeping it constant throughout the entire preparation process; slowly adding 200 g of an active ingredient, such as mannitol; cooling the solution until it reaches a temperature of less than 35°C; adding 1.0 g of sodium phosphate monobasic monohydrate; adding 5.1 g of sodium phosphate dibasic anhydrous; adding 1.0 ml of Sophisen; bringing it to a volume of 1 liter

with injectable water; and agitating at 200 rpm ± 50 rpm until complete homogeneity is obtained.

16. An intravitreous injectable solution for the treatment of vitreous hemorrhages comprising: a pharmaceutically effective quantity of mannitol: 0.01% to 5% in weight of sodium phosphate monobasic monohydrate; 0.01% to 5% in weight of sodium phosphate dibasic anhydrous and 100 ml of injectable water.

17. The intravitreous injectable solution of claim 16, in which the mannitol is present in a percentage of 5% to 30% in weight of the solution.

18. The intravitreous injectable solution of claim 16, which further includes 0.05% to 20% in weight of a carrier solution.

19. The intravitreous injectable solution of claim 16, wherein the carrier solution is Sophisen.